

IDS Form PTO/SB/08: Substitute for form 1449A/P

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# INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(Use as many sheets as necessary)

Application Number	10/573,352
Filing Date	March 24, 2006
First Named Inventor	Robert Hugh Bradbury
Art Unit	1624
Examiner Name	TRUONG, Tamthom
Attorney Docket Number	09953.0010-00000

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## U.S. PATENTS AND PUBLISHED U.S. PATENT APPLICATIONS

Examiner Initials	Cite No.	Document Number	Issue or Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code (if known)			
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Examiner Signature	/Tamthom Truong/	Date Considered	02/27/2009
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IDS Form PTO/SB/08: Substitute for form 1449A/PTO				<b>Complete if Known</b>	
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Sheet	2	of	10	Attorney Docket Number	09963 0010-00000

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Examiner Initials	Cite No.	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	Translation
		Country Code Number Kind Code (if known)				
		WO 2007/063293	06-07-2007	Astrazeneca UK Ltd		

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Examiner Name	TRUONG, Tamthom
Attorney Docket Number	09953.0010-00000

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## **NON PATENT LITERATURE DOCUMENTS**

Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	Translation
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IDS Form PTO/SB/08; Substitute for form 1449A/PTO				<b>Complete if Known</b>	
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  <i>(Use as many sheets as necessary)</i>				Application Number	10/573,352
				Filing Date	March 24, 2006
				First Named Inventor	Robert Hugh Bradbury
				Art Unit	1624
				Examiner Name	TRUONG, Tamthom
Sheet	9	of	10	Attorney Docket Number	09963.0010-00000

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	Translation
		Myers et al., "The preparation and SAR of 4-(anilino), 4-(phenoxy), and 4-(thiophenoxy)-quinazolines: inhibitors of p56lck and EGF-R tyrosine kinase activity," <i>Biorg. Med. Chem. Lett.</i> 7(4): 417-420 (1997).	
		Pao et al., "Epidermal Growth Factor Receptor Mutations, Small-Molecule Kinase Inhibitors, and Non-Small-Cell Lung Cancer: Current Knowledge and Future Directions," <i>Journal of Clinical Oncology</i> 23(11):1-13 (2005).	
		Rewcastle et al., "Tyrosine Kinase Inhibitors. 5 ... 4-(Phenylamino)quinazolines as Potent ... Inhibitors of the Tyrosine Kinase Domain of the Epidermal Growth Factor Receptor," <i>J. Med. Chem.</i> 38:3482-3487 (1995).	
		Singh et al., "Inhibitors of the epidermal growth factor receptor protein tyrosine kinase: A quantitative structure-activity relationship analysis," <i>J. Enzyme Inhibition</i> 13:125-134 (1998).	
		Small et al., "Tyrosine kinase inhibitors. 17. Irreversible inhibitors of the epidermal growth factor receptor: 4-(Phenylamino)quinazoline- and 4-(Phe-nylamino)pyrido," <i>J. Med. Chem.</i> 43(16):3199 (2000).	
		Stamos et al., "Structure of the Epidermal Growth Factor Receptor Kinase Domain Alone and in Complex with a 4-Anilinoquinazoline Inhibitor," <i>J. Bio. Chem.</i> 277(48):46265-46272 (2002).	
		Traxler, "Oncologic, Endocrine & Metabolic: Protein tyrosine kinase inhibitors in cancer treatment," <i>Expert Opinion on Therapeutic Patents</i> 7:571-588 (1997).	
		Traxler, "Monthly Focus: Oncologic, Endocrine & Metabolic: Tyrosine kinase inhibitors in cancer treatment (Part II)," <i>Expert Opinion on Therapeutic Patents</i> 8:1599-1625 (1998).	
		Tsou et al., "6- Substituted-4-(3-bromophenylamino)quinazolines As Putative Irreversible Inhibitors of the Epidermal Growth Factor Receptor (Egfr) and Human Epidermal Growth Factor Receptor (Her-2) Tyrosine Kinases with Enhanced Antitumor Activity," <i>J. Med. Chem.</i> 44:2719-2734 (2001).	
		Vema et al., "Design of EGFR Kinase Inhibitors: A Ligand-Based Approach and Its Confirmation with Structure-Based Studies," <i>Bioorg. Med. Chem.</i> 11:4643-4653 (2003).	
		Wright et al., "Allosteric inhibition of fructose-1,6-bisphosphatase by anilinoquinazolines," <i>Bioorg. Med. Chem. Lett.</i> 11(1):17-21 (2001).	
		Decision in Patent Interferences 105,595 McK and 105,596 McK dated June 17, 2008.	
		English Translation of Office Action in Japanese Patent Appln. No. 2003-580299, the Japanese counterpart of the present application, dated May 11, 2006.	
		Response to Office Action in Japanese Patent Appln. No. 2003-580299, the Japanese counterpart of the present application, dated July 28, 2006.	
		English translation of Response to Office Action in Japanese Patent Appln. No. 2003-580299, the Japanese counterpart of the present application, dated July 28, 2006.	

Examiner Signature	/s/ Tamthom Truong/	Date Considered	02/27/2009
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		English translation of Response to Office Action in Japanese Patent Appln. No. 2003-580299, the Japanese counterpart of U.S. Application No. 10/508,675, dated October 26, 2006.	
		Office Action in Indian Patent Appln. No. 2630/DELNP/2004, the Indian counterpart of the present application, dated April 20, 2006.	
		Response to Office Action in Indian Patent Appln. No. 2630/DELNP/2004, the Indian counterpart of the present application, dated July 24, 2006.	
		English translation of Office Action in Chinese Patent Appln. No. 03811739.8, the Chinese counterpart of the present application, dated July 21, 2006.	
		Response to Office Action in Chinese Patent Appln. No. 03811739.8, the Chinese counterpart of U.S. Application No. 10/508,675, dated December 5, 2006.	
		English translation of Response to Office Action in Chinese Patent Appln. No. 03811739.8 of U.S. Application No. 10/508,675, dated December 5, 2006.	
		Communication from European Patent Office in EP Appln. No. 03 710 015.3, the European counterpart of the present application, dated September 22, 2006.	
		Communication from EPO dated March 9, 2006, in EP Appln. No. 03 710 015.3, the European counterpart of U.S. Application No. 10/508,675	
		Communication from European Patent Office ("EPO") dated May 27, 2005, in EP Appln. No. 03 710 015.3, the European counterpart of U.S. Application No. 10/508,675	
		Reply to May 27, 2005 Communication from EPO dated September 20, 2005, in EP Appln. No. 03 710 015.3, the European counterpart of U.S. Application No. 10/508,675	
		Office Action in copending US Application No. 10/571,991 mailed August 19, 2008.	
		G.A. Patani et al., "Bioisosterism: A Rational Approach in Drug Design," <i>Chem. Rev.</i> 96:3147-3176 (1996).	
		Harris et al, Poster presented at the XXII European Colloquium on Heterocyclic Chemistry (XXII ECHC-2006) in Bari, Italy, September 2 - 6, 2006.	

Examiner Signature	/Tamthom Truong/	Date Considered	02/27/2009
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